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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/605,293	06/28/2000	DAVID L. CHAPEK	MIO-0037-VA	5927	
7590 03/03/2004  KILLWORTH GOTTMAN HAGAN SCHAEFF L L P ONE DAYTON CENTRE, SUITE 500			EXAMINER		
			RICHARDS, N DREW		
DAYTON, OH 45402-2023			ART UNIT	PAPER NUMBER	
			2815		

DATE MAILED: 03/03/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

			1 <sup>v</sup> .
	Application No.	Applicant(s)	
Office Astis Comment	09/605,293	CHAPEK, DAVID L.	
Office Action Summary	Examiner	Art Unit	
	N. Drew Richards	2815	
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with the	correspondence address	
A SHORTENED STATUTORY PERIOD FOR REPL THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a repleved in the period for reply is specified above, the maximum statutory period Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	136(a). In no event, however, may a reply be to bly within the statutory minimum of thirty (30) da will apply and will expire SIX (6) MONTHS from e, cause the application to become ABANDON	imely filed  ys will be considered timely.  n the mailing date of this communication.  ED (35 U.S.C. § 133).	
Status			
1)⊠ Responsive to communication(s) filed on <u>05 L</u>	December 2003.		
	s action is non-final.		
3) Since this application is in condition for allows closed in accordance with the practice under	·		
Disposition of Claims			
4) ☐ Claim(s) 9-12 and 14 is/are pending in the ap 4a) Of the above claim(s) is/are withdra 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 9-12 and 14 is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or	awn from consideration.		
Application Papers			
9)☐ The specification is objected to by the Examin	er.		
10)⊠ The drawing(s) filed on <u>28 June 2000</u> is/are: a			
Applicant may not request that any objection to the	÷, ,	` *	
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the E	• • • • • • • • • • • • • • • • • • • •	•	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureat * See the attached detailed Office action for a list	ts have been received. ts have been received in Applica prity documents have been receiv nu (PCT Rule 17.2(a)).	tion No ved in this National Stage	
Attachment(s)			
1) Notice of References Cited (PTO-892)	4) Interview Summar		
<ul> <li>Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08 Paper No(s)/Mail Date</li> </ul>	Paper No(s)/Mail [ 5) Notice of Informal 6) Other:	Patent Application (PTO-152)	

#### **DETAILED ACTION**

## Claim Rejections - 35 USC § 103

- 1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
  - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 2. Claim 9 is rejected under 35 U.S.C. 103(a) as being unpatentable over Applicant's admitted prior art in view of Henley et al. (U.S. Patent No. 6083324).

Applicant's admitted prior art discloses on page 1 lines 12-16 a semiconductor substrate, a layer of silicon dioxide on the substrate, and a layer of polycrystalline silicon formed on the silicon dioxide, the polycrystalline silicon having a smooth morphology. The admitted prior art discloses the layer of silicon dioxide having been doped with hydrogen ions. The semiconductor substrate is considered as a bottom portion of the silicon dioxide layer with the remaining silicon dioxide layer as the silicon dioxide layer upon the substrate. Though the admitted prior art does not explicitly state a layer of polysilicon is on the silicon dioxide it is implicitly understood that the polysilicon is formed seeing that the admitted prior art discusses performing the hydrogen doping of the silicon dioxide so as to provide a thinner, smoother polysilicon film deposited on the silicon dioxide. The admitted prior art of lines 16-22 does not teach the layer of silicon dioxide being free of sputtered metal contaminants as the Kaufman ion source causes metal contaminants in the layer. The admitted prior art teaches the metal contaminants being produced from metal sputtering off a metal grid in the Kaufman ion source

apparatus and that as device sizes decrease the effect of the damage from the metal contaminants increases.

Henley et al. teach a plasma immersion ion implantation apparatus for implanting hydrogen ions into semiconductors. It is noted that plasma source ion implantation (PSII or PSI) and plasma immersion ion implantation (PIII) are interchangeable terms for the same plasma treatment. Henley et al. teach that their implantation method can be used on SOI (silicon-on-insulator) substrates in column 2 lines 30-40 and teach implanting hydrogen ions in line 38, for example.

Applicants admitted prior art and Henley et al. are combinable because they are from the same field of endeavor. At the time of the invention it would have been obvious to a person of ordinary skill in the art to implant the hydrogen using the PSII technique of Henley et al. The motivation for doing so is to that PIII is cost effective, easy to use, and produces less impurity metal contamination. Therefore, it would have been obvious to combine Applicant's admitted prior art with Henley et al. to obtain the invention of claim 9.

3. Claims 10-12 are rejected under 35 U.S.C. 103(a) as being unpatentable over Burns et al. (Principles of Electronic Circuits, Pp. 380 and 381) in view of Applicant's admitted prior art with Henley et al. as applied to claim 9 above.

Burns et al. teach a field effect transistor in figure 9.8 on page 381. Burns et al. teach a substrate, a silicon dioxide layer formed on at least a portion of the substrate, a layer of polycrystalline silicon formed on at least a portion of the silicon dioxide layer,

and a gate oxide formed on the substrate from the layer of silicon dioxide, and a source and a drain in the substrate where a gate electrode is formed on the substrate from the layer of polycrystalline silicon. Burns et al. do not teach the layer of silicon dioxide having hydrogen ions implanted therein or being free of sputtered metal contaminants. Applicant's admitted prior art teaches implanting hydrogen ions into silicon dioxide on page 1 lines 12-16 and Henley et al. teach using a PSII method to implant the hydrogen. Applicant's admitted prior art with Henley et al. as discussed above also teach the silicon dioxide as being free of sputtered metal contaminants. In the combination of the references, the gate oxide would be formed from the layer of silicon dioxide having hydrogen ions implanted therein and the layer of polycrystalline silicon formed on the layer of silicon dioxide would have a smooth morphology.

Burns et al. and Applicant's admitted prior art with Henley et al. are combinable because they are from the same field of endeavor. At the time of the invention it would have been obvious to a person of ordinary skill in the art to implant hydrogen ions into the silicon dioxide layer. The motivation for doing so is to prepare the surface of the silicon dioxide for the deposition of a layer of polycrystalline silicon to provide for a thinner and smoother polycrystalline silicon film. Therefore, it would have been obvious to combine Burns et al. with Applicant's admitted prior art and Henley et al. to obtain the invention of claim 10.

With regard to claim 11, Burns et al. teach on pages 380 and 381, a memory array which further includes a plurality of memory cells arranged in rows and columns

comprising at least one field effect transistor having a gate oxide, source, and drain formed on the substrate and a gate electrode for each transistor formed of the layer of polycrystalline silicon. The gate oxide for each transistor of the combination of references would be formed of the silicon dioxide having hydrogen atoms implanted therein.

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With regard to claim 12, Official Notice is taken that one of ordinary skill in the art at the time of the invention would form the transistor of claim 10 or the memory array of claim 11 on a semiconductor wafer including a plurality of die. This is well known as in semiconductor processing multiple devices are formed on a single wafer then split into individual die to allow for processing of a great number of die at one time to save of processing costs. Also, the gate electrode is a repeating series of gate electrodes for each transistor on each die formed from the layer of polycrystalline silicon.

4. Claim 14 is rejected under 35 U.S.C. 103(a) as being unpatentable over Murata et al. (U.S. Patent No. 5576229) in view of Applicant's admitted prior art with Henley et al. as applied to claims 9-12 above.

Murata et al. teach a thin film transistor in figure 6E comprising a semiconductor substrate 501 of glass, a layer of polycrystalline silicon 507 formed on a portion of the substrate, a insulating layer 503 formed on a portion of the polycrystalline silicon, a gate oxide formed from the insulating layer, a source region 507a and drain region 507b formed in the polycrystalline silicon, and a gate electrode 504 formed on the insulating

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layer. Murata et al. do not teach the substrate having hydrogen ions implanted therein or the substrate being free of sputtered metal contaminants. Applicant's admitted prior art teaches implanting hydrogen ions into a silicon dioxide (glass) layer to provide a smooth topology polycrystalline silicon film thereon on page 1 lines 12-16. Henley et al. as discussed above teach using a PSII method to implant the hydrogen. Applicant's admitted prior art with Henley et al. as discussed above also teach the silicon dioxide as being free of sputtered metal contaminants. In the combination of the references, the layer of polycrystalline silicon formed on the layer of substrate would have a smooth morphology.

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Murata et al. and Applicant's admitted prior art with Henley et al. are combinable because they are from the same field of endeavor. At the time of the invention it would have been obvious to a person of ordinary skill in the art to implant hydrogen ions into the glass substrate. The motivation for doing so is to prepare the surface of the glass substrate for the deposition of a layer of polycrystalline silicon to provide for a thinner and smoother polycrystalline silicon film. Therefore, it would have been obvious to combine Murata et al. with Applicant's admitted prior art and Henley et al. to obtain the invention of claim 14.

#### Response to Arguments

5. Applicant's arguments filed 12/5/03 have been fully considered but they are not persuasive.

First, applicant has argued that Henley et al. provide no motivation to pick hydrogen from the numerous possible choices of ions that Henley et al. teach may be implanted. This is not persuasive as Henley et al. is not relied upon for the teaching of choosing hydrogen to implant. Hydrogen implantation is taught by the admitted prior art. Henley et al. is merely relied upon to teach that hydrogen may be implanted into semiconductor articles by a PSII method as claimed.

Second, applicant has argued that there is no motivation to combine the teachings of the admitted prior art with Henley et al. This is not persuasive as the admitted prior art clearly acknowledges the problems associated with metal contamination in the implantation of hydrogen into silicon dioxide (see page 1 lines 18-20 of the applicant's specification). Henley et al. explicitly states that the PIII (same method as PSII) method produces less impurity metal contamination than other ion implantation methods (see column 3 lines 13-17 of Henley et al.). In light of the admitted prior art's need to reduce metal contamination one of ordinary skill in the art would have been motivated to use the ion implantation method of Henley et al. because of the stated advantage of reducing the problematic metal contamination.

Applicant also argues that Henley et al. does not teach implanting into a silicon dioxide substrate or implanting into a silicon dioxide substrate for the purpose of providing a subsequent layer of polycrystalline silicon which has a smooth surface.

Henley et al. was not relied upon to teach implanting into a silicon dioxide substrate or doing so for the stated purpose. These limitations were taught by the admitted prior art.

Applicant also argues that Henley et al. teach away from treating the insulating layer by specifically implanting ions above the insulating layer and thus does not teach a surface treatment. Again, Henley et al. was not relied upon to teach implanting ions into the insulating layer or for a surface treatment, these limitations were taught by the admitted prior art as explained in the rejections above. Henley et al. was relied upon to correct the deficiencies of the admitted prior art, i.e. metal contamination from the Kaufman ion source. Henley et al. explicitly teaches that hydrogen ions can be implanted into semiconductor articles by PIII and that the PIII produces less impurity metal contamination. Thus, Henley et al. teaches that PIII can be used for a process similar to that claimed (implanting hydrogen ions) and provides proper motivation for why one would use the PIII method. Therefore, the arguments are not persuasive and the rejections are considered proper.

## Conclusion

6. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of

the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to N. Drew Richards whose telephone number is (571) 272-1736. The examiner can normally be reached on M-F 8:00-5:30; Every other Friday off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Tom Thomas can be reached on (571) 272-1664. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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